

ABSTRACT

The present invention relates to a rapidly disintegrable tablet for oral administration, which disintegrates in the oral cavity within 60 seconds, consisting essentially of (i) a therapeutically effective amount of an active ingredient, (ii) spray-dried mannitol, of which at least 80% has an average particle size over 100 μm , (iii) crospovidone, and (iv) one or more pharmaceutically acceptable excipients, the tablet containing no microcrystalline cellulose.

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